SEARCH NOTES 10 735, 439

Royds 10/735,439

04/26/2006

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> d his nofil
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(FILE 'HOME' ENTERED AT 09:39:49 ON 26 APR 2006)
    FILE 'REGISTRY' ENTERED AT 09:40:12 ON 26 APR 2006
               E 3,3,14,14-TETRAMETHYL/CN
L1
              1 SEA ABB=ON PLU=ON "3,3,14,14-TETRAMETHYLHEXADECANEDIOIC
               ACID"/CN
               D SCA
               D
               STR 87272-20-6
L2
L3
             0 SEA FAM SAM L2
             1 SEA FAM FUL L2
L4
    FILE 'HCAPLUS' ENTERED AT 09:42:15 ON 26 APR 2006
L5
     43 SEA ABB=ON PLU=ON L4
    FILE 'MEDLINE, EMBASE, BIOSIS' ENTERED AT 09:43:02 ON 26 APR 2006
L6
            65 SEA ABB=ON PLU=ON L4
     FILE 'HCAPLUS' ENTERED AT 09:43:12 ON 26 APR 2006
               E SYNDROME X/CT
               E E3+ALL
               E E2+ALL
L7
          2863 SEA ABB=ON PLU=ON "DISEASE, ANIMAL (L) METABOLIC SYNDROME
               X"+PFT, NT/CT
               E DYSLIPOPROTEINEM/CT
               E E4+ALL
               E E2+ALL
           227 SEA ABB=ON PLU=ON "LIPOPROTEINS (L) DYSLIPOPROTEINEMIA"+PFT, N
rs
               T/CT
               E PLASMA TRIGLYCERIDES/CT
               E E3+ALL
               E E2+ALL
         20505 SEA ABB=ON PLU=ON "GLYCERIDES (L) BLOOD"+PFT, NT/CT
L9
               E HDL CHOLESTEROL/CT
               E HDL/CT
               E E3+ALL
               E CHOLESTEROL/CT
               E E3+ALL
               E HDL/CT
               E E3+ALL
               E E2+ALL
         24207 SEA ABB=ON PLU=ON "LIPOPROTEINS (L) HIGH-D."+PFT,NT/CT
L10
L11
        208295 SEA ABB=ON PLU=ON (L7 OR L8 OR L9 OR L10) OR SYNDROM? (2A) X
               OR TRIGLYCERID? OR HDL OR CHOLESTEROL? OR DYSLIPOPROT?
L12
            19 SEA ABB=ON PLU=ON L4 AND L11
     FILE 'MEDLINE' ENTERED AT 09:49:56 ON 26 APR 2006
L13
            28 SEA ABB=ON PLU=ON L4
               E SYNDROME X/CT
     FILE 'MEDLINE, EMBASE, BIOSIS' ENTERED AT 09:50:33 ON 26 APR 2006
L14
            65 SEA ABB=ON PLU=ON L4
        528315 SEA ABB=ON PLU=ON SYNDROM? (2A) X OR TRIGLYCERID? OR HDL OR
L15.
               CHOLESTEROL? OR DYSLIPOPROT? OR HIGH(3A) (D OR DENS?) (3A) (LIP?
               OR CHOLESTER?)
L16
            27 SEA ABB=ON PLU=ON L14 AND L15
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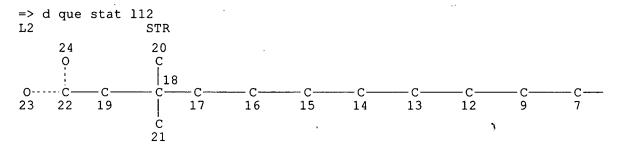
=> fil hcap FILE 'HCAPLUS' ENTERED AT 09:52:42 ON 26 APR 2006 USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT. PLEASE SEE "HELP USAGETERMS" FOR DETAILS. COPYRIGHT (C) 2006 AMERICAN CHEMICAL SOCIETY (ACS)

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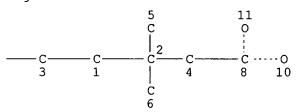
FILE COVERS 1907 - 26 Apr 2006 VOL 144 ISS 18 FILE LAST UPDATED: 25 Apr 2006 (20060425/ED)

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Page 1-A



Page 1-B NODE ATTRIBUTES: DEFAULT MLEVEL IS ATOM DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES: RING(S) ARE ISOLATED OR EMBEDDED NUMBER OF NODES IS 24

STEREO ATTRIBUTES: NONE

L4 1 SEA FILE=REGISTRY FAM FUL L2

L7 2863 SEA FILE=HCAPLUS ABB=ON PLU=ON "DISEASE, ANIMAL (L) METABOLIC SYNDROME X"+PFT, NT/CT

r8	227		PLU=ON	"LIPOPROTEINS (L) DYSLIPOPROTE
		INEMIA"+PFT,NT/CT		
L9	20505		PLU=ON	"GLYCERIDES (L) BLOOD"+PFT,NT/
		CT		
L10	24207		PLU=ON	"LIPOPROTEINS (L) HIGH-D."+PFT
		,NT/CT	,	
L11	208295			(L7 OR L8 OR L9 OR L10) OR
		SYNDROM? (2A) X OR TRIGLY	CERID? OR	HDL OR CHOLESTEROL? OR
		DYSLIPOPROT?		
L12	19	SEA FILE=HCAPLUS ABB=ON	PLU=ON	L4 AND L11

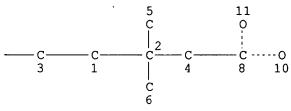
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FILE 'BIOSIS' ENTERED AT 09:52:55 ON 26 APR 2006 Copyright (c) 2006 The Thomson Corporation

=> d que stat 116 L2 STR 20 24 0 С 18 O----- C-- C-19 17 13 12 23 22 16. 15 14 9 Ċ 21





Page 1-B NODE ATTRIBUTES: DEFAULT MLEVEL IS ATOM DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES: RING(S) ARE ISOLATED OR EMBEDDED NUMBER OF NODES IS 24

STEREO ATTRIBUTES: NONE

L4 1 SEA FILE=REGISTRY FAM FUL L2

L14 65 SEA L4

L15 528315 SEA SYNDROM? (2A) X OR TRIGLYCERID? OR HDL OR CHOLESTEROL? OR DYSLIPOPROT? OR HIGH (3A) (D OR DENS?) (3A) (LIP? OR CHOLESTER?)

L16 27 SEA L14 AND L15

=> dup rem 112 116 FILE 'HCAPLUS' ENTERED AT 09:53:03 ON 26 APR 2006 USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT. PLEASE SEE "HELP USAGETERMS" FOR DETAILS. COPYRIGHT (C) 2006 AMERICAN CHEMICAL SOCIETY (ACS)

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FILE 'BIOSIS' ENTERED AT 09:53:03 ON 26 APR 2006 Copyright (c) 2006 The Thomson Corporation PROCESSING COMPLETED FOR L12 PROCESSING COMPLETED. FOR L16 L17 26 DUP REM L12 L16 (20 DUPLICATES REMOVED)

ANSWERS '1-19' FROM FILE HCAPLUS ANSWERS '20-21' FROM FILE MEDLINE ANSWERS '22-26' FROM FILE EMBASE

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L17 ANSWER 1 OF 26 HCAPLUS COPYRIGHT 2006 ACS on STN DUPLICATE 1

ACCESSION NUMBER: 2005:1162213 HCAPLUS

DOCUMENT NUMBER: 144:408

SOURCE:

TITLE: Transcriptional suppression of human microsomal

triglyceride transfer protein by hypolipidemic

insulin sensitizers

AUTHOR(S): Sheena, Vered; Hertz, Rachel; Berman, Ina; Nousbeck,

Janna; Bar-Tana, Jacob

Department of Human Nutrition and Metabolism, Hebrew CORPORATE SOURCE:

University Medical School, Jerusalem, 91120, Israel Biochemical Pharmacology (2005), 70(11), 1548-1559

CODEN: BCPCA6; ISSN: 0006-2952

PUBLISHER: Elsevier B.V. DOCUMENT TYPE: Journal LANGUAGE: English

Microsomal triglyceride transfer protein (MTP) catalyzes the assembly and secretion of liver triglyceride-rich lipoproteins. The human MTP (hMTP) promoter activity is reported here to be suppressed by $HNF-4\alpha$ ligand antagonists (e.g., Medica analogs) or by PPARy ligand agonists (e.g., thiazolidinediones), thus accounting for their hypolipidemic activity in humans. Suppression of liver hMTP by Medica analogs or by thiazolidinediones was mediated by the TAAA sequence that serves as non-canonical TATA box of the hMTP core promoter. MTP suppression was evident in the specific context of the wild type hMTP core promoter, but not in the context of the mutated rodent-conforming hMTP core promoter governed by a canonical TATA box conjoined with its proximal (-50/-38) DR-1 element. HMTP suppression by Medica analogs or thiazolidinediones mediated by hMTP TAAA was independent of ${\tt HNF-4}\alpha$ or PPARy. HMTP suppression by Medica analogs, but not by thiazolidinediones, was further complemented by inhibition of $HNF-4\alpha$ transcriptional activity transduced by the distal (-83/-70) DR-1 element of hMTP promoter. HMTP promoter activity was unaffected by PPARa activation. Furthermore, in contrast to hMTP, the promoter activity of the rodent-conforming hMTP was robustly activated by Wy-14,643-activated PPARα or by thiazolidinedione-activated PPARγ. Transcriptional activation by PPARa or PPARy of the rodent-conforming, but not the wild type hMTP gene promoter, resulted from => d his nofil

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(FILE 'HOME' ENTERED AT 11:25:02 ON 26 APR 2006)
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FILE 'HCAPLUS' ENTERED AT 11:25:14 ON 26 APR 2006

E BAR-TANA/AU E BAR TANA/AU

105 SEA ABB=ON PLU=ON ("BAR TANA J"/AU OR "BAR TANA JACOB"/AU) L1E BARTANA/AU

FILE 'REGISTRY' ENTERED AT 11:26:23 ON 26 APR 2006

E 3,3,14,14/CN

E 3,3,14,14-TETRAMETHYL/CN

1 SEA ABB=ON PLU=ON "3,3,14,14-TETRAMETHYLHEXADECANEDIOIC L2 ACID"/CN

FILE 'HCAPLUS' ENTERED AT 11:26:57 ON 26 APR 2006

L3

35 SEA ABB=ON PLU=ON L1 AND L2 20 SEA ABB=ON PLU=ON L1 AND (SYNDROM?(2A)X OR DYSLIPOPROT? OR L4

HDL OR CHOLESTEROL? OR TRIGLYCERID?)

15 SEA ABB=ON PLU=ON L3 AND L4 L5 40 SEA ABB=ON PLU=ON L3 OR L4 L6

 \Rightarrow d 16 ibib abs hitind 1-40

ANSWER 1 OF 40 HCAPLUS COPYRIGHT 2006 ACS on STN

2005:1162213 HCAPLUS ACCESSION NUMBER:

DOCUMENT NUMBER: 144:408

TITLE: Transcriptional suppression of human microsomal

triglyceride transfer protein by hypolipidemic

insulin sensitizers

AUTHOR(S):Sheena, Vered; Hertz, Rachel; Berman, Ina; Nousbeck,

Janna; Bar-Tana, Jacob

Department of Human Nutrition and Metabolism, Hebrew CORPORATE SOURCE:

University Medical School, Jerusalem, 91120, Israel

Biochemical Pharmacology (2005), 70(11), 1548-1559 SOURCE:

CODEN: BCPCA6; ISSN: 0006-2952

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